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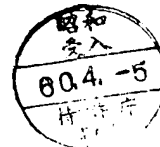
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Bloomsbury Way, London WC1A 2TP (GB)(54) **The use of dispersing aids in processing aid compositions for vinyl chloride polymer, the production of said compositions and vinyl chloride compositions containing them.**(57) The dispersability of processing aids for vinyl chloride polymers is improved by the use of a dispersing aid comprising at least one homopolymer and/or copolymer containing at least 50% by weight of units of one or more of the following monomers: (C₁-C₈)-alkyl acrylates, (C₁-C₉)-cycloalkyl acrylates, ethylene, propylene, vinyl acetate and butadiene.Preferred processing aid compositions therefor comprise blends of (a) copolymer(s) of methyl methacrylate and, in an amount of less than 50% by weight of the copolymer, (C₁-C₄)-alkyl acrylate(s) and (b) at least one such dispersing aid homopolymer or copolymer wherein the ratio of (a):(b) is 99.5:0.5 to 90:10. The blend may be prepared by intimately mixing components (a) and (b) as latex polymer and subsequently coisolated.**EP 0 132 317 A1**

THE USE OF DISPERSING AIDS IN PROCESSING AID COMPOSITIONS
FOR VINYL CHLORIDE POLYMER, THE PRODUCTION OF SAID
COMPOSITIONS AND VINYL CHLORIDE COMPOSITIONS CONTAINING
THEM.

5 This invention is concerned with the use of
dispersing aids in processing aid compositions for vinyl
chloride polymer, the production of said compositions and
vinyl chloride compositions containing them.

10 Polymeric processing aids are widely used in the
production of poly(vinyl chloride) articles, in order to
improve the processing performance during extrusion,
injection and/or thermoforming. The importance of
dispersing the processing aids well through the vinyl
15 chloride polymer is known. Improvements in dispersion of
such processing aids in poly(vinyl chloride) have been
achieved by the use of high friction conditions or
temperature variation during processing. Also, it is known
that lumps of processing aid polymer particles which do not
20 break down during processing can form gels in the final
product and these are visible and detract from optimum
processability.

 In spite of the wide recognition of this dispersion
problem, no adequate generally usable solution has been
available. We have now found certain dispersion aid
25 compositions which, when used to modify the processing aids,
have a reduced tendency to form gel colonies and so have
desirable dispersability properties. In its broadest aspect
this invention therefore is directed to this use.

 Processing aid compositions of this invention
30 preferably comprise blends of (a) copolymer(s) of methyl
methacrylate with less than 50% by weight of (C₁-C₄)-alkyl
acrylate(s) and (b) homopolymer(s) or copolymer(s)
containing at least 50% by weight of one or more of the
monomers:

35 (C₁ to C₈-alkyl) acrylates, (C₆ to C₉)-cycloalkyl acrylates,
ethylene, propylene, vinyl acetate, butadiene; in a weight
ratio (a):(b) of 99.5:0.5 to 90:10 preferably 99:1 to 95:5.

Consequently the invention also provides compositions comprising vinyl chloride polymer containing the processing aid compositions of the invention and processes for making the processing aid compositions by intimately mixing components (a) and (b) in the indicated weight ratios.

The processing aid polymers (a) which are used in this invention may be single stage polymers, or, multiple stage polymers, and the Examples of such processing aids are more fully described in British Patents 1 511 683, 981 116 and U.S. Patent 3 833 686.

The component (b), is preferably at least one of the following polymers:

poly(butyl acrylate), poly(ethyl acrylate), polybutadiene, poly(ethyl acrylate/styrene), poly(butyl acrylate/styrene), poly(butyl acrylate/methyl methacrylate), poly(ethyl acrylate/butyl methacrylate), poly(butadiene/styrene), poly(butyl acrylate/styrene/methylmethacrylate) with the proviso that the methacrylate and/or styrene content is less than 50% by weight in each case. Other examples are poly(vinyl acetate) and poly(ethylene/vinyl acetate).

The molecular weight range of suitable dispersing aids is wide, for example 15,000 to 100,000 weight average, but depends on the particular polymer used. For example, with poly(butyl acrylate), poly(ethyl acrylate), and poly(butyl acrylate/styrene), a preferred molecular weight range is as stated above. The dispersing aid may even be cross-linked and/or graft linked preferably to a maximum content of 5% by weight crosslinker or graft-linker units. A dispersability rating test for indicating the suitability of a particular polymer for use as a dispersing aid is given later in this specification.

The dispersing aid is intimately blended with the processing aid polymer to form the compositions of the invention. Any method of making a very thorough, intimate mixture of the dispersing aid and usually powder processing aid polymer is suitable. However simple admixture of

processing aid powder and dispersing aid will usually be insufficient unless intimate, thorough mixing is achieved. Another method of forming the compositions of the invention is to emulsion polymerise the dispersing aids in the presence of the latex of the processing aid polymer formed by emulsion polymerisation. Our presently preferred method is to separately prepare emulsions of processing aid polymer and dispersing aid, mix the emulsions, and then coisolate the two polymers by spray drying or coagulation. Spray drying is described in detail in Spray Drying, An Introduction to Principles, Operational Practices, and Applications, K. Masters, CRC Press, Cleveland, Ohio (1972). Coagulation comprises salting out the stabilizing surface active agent from the emulsion by slowly adding the polymer emulsion for example twice its volume of electrolyte solution containing any one or more of a number of salts and/or acids such as NaCl, KCl, CaCl₂, Na₂SO₄, MgSO₄, acetic acid, oxalic acid, HCl, HNO₃, H₂SO₄ and H₃PO₄.

The polymer may then be separated by filtration and washed several times with fresh water to remove excess electrolyte, and then dried.

The processing aid and dispersing aid can also be isolated from the emulsion by evaporation or freeze drying. When following the preferred method, it is important that the two emulsions be compatible. For instance, the soaps should be similar or the same, i.e., of the same ionic species and of similar water solubility or HLB (hydrophile/lipophile balance).

The dispersing aid is generally a single stage polymer but it may be a multiple stage polymer in which case the overall monomer composition and the outermost (shell) stage must both conform to the composition parameters specified above even if some of the early (inner) stages, for example the core, do not. The dispersing aid may intentionally be separately prepared as distinct particles and then blended

with the processing aid polymer. The dispersing aid polymer is not to be confused with any ungrafted polymer which unintentionally forms during preparation of multiple stage processing aid polymers, since such unintentional polymer does not function to reduce gel formation and is thus not within the scope of this invention.

The most preferred weight ratio of processing aid polymer to dispersing aid in the compositions of the invention is 98/2 to about 96/4, and the optimum weight ratio is about 97/3.

The vinyl chloride polymers which may be modified using the compositions of the invention include polymers and copolymers of vinyl chloride containing at most 50% by weight of other monomer units, for example vinyl acetate, ethyl vinyl acetate.

The weight ratio of vinyl chloride polymer to processing aid composition may be the same as used with conventional processing aids. Normally this ratio is 90:10 to 99;5:0.5 more usually 95:5 to 99:1 by weight.

Other standard additives may be used to make up the formulation of the vinyl chloride polymer composition. For example, impact modifiers, colorants, pigments, plasticizers, stabilizers, and lubricants may be included.

After formulating by standard techniques, the formulation may be melt processed, also by standard techniques, to form a substantially gel-free compound in which the processing aid is extremely well dispersed.

Some preferred embodiments of the invention are given in the following Examples. As comparative tests there are also given details of the performance of conventional processing aids and details of tests in which the candidate as dispersing aid fails the dispersability rating test previously mentioned and detailed below. All parts and percentages are by weight unless otherwise indicated.

Candidate dispersing aid polymers of varying compositions are prepared in emulsion using the following

ingredients and procedure.

<u>Charges:</u>		<u>Parts:</u>
	A - DI water @ pH 10 using NaOH	397.5
	B - sodium stearate (or SLS)	0.26
5	B1 - sodium stearate (or SLS)	1.49
	monomer(s)	212.6
	C - t-dodecyl mercaptan as mercaptan chain transfer agent or BDA as	
10	difunctional monomer	Varied
	D - KPS initiator	1.1
	water	63.3
	Theory Solids	<u>32.4%</u>

Procedure

15 Charge the water (A) to a one litre four-neck flask fitted with a stirrer, thermometer, nitrogen inlet and a condenser. The water is heated to 65°C while sweeping with nitrogen. At 65°C the first emulsifier charge (B) is added and allowed to stir for 10 minutes. The KPS initiator (D)

20 is added followed by 15% of the monomer(s) (C) and the temperatures maintained at 65°C until an exotherm is observed.

25 At this point the second emulsifier charge (B1) is added followed by the gradual addition of the balance of the monomers (C) over a one hour period. After completion of the monomer feed, the batch is maintained at 65°C for 15 minutes then cooled to room temperature.

A PVC masterbatch formulation, without dispersing aid, of the following ingredients

	PVC (K-58) Resin	100 Parts
5	Sulfur containing organic tin stabilizer	2.0 Parts
	Stearic Acid	1.0 Part
	Plasticizer-DOP	5.0 Parts
	Lubricating-Processing Aid	
	"Paraloid" K-175	3.0 Parts

10 is prepared in a high intensity Prodex-Henschel blender. The subject processing aid composition powders are added (at 108g masterbatch + 2g processing aid) to samples of the masterbatch and the powder is again mixed homogenously to form the final compound.

15 The powder compound is milled on a two-roll Schwabenthan Mill having two chrome polished rolls. The dimensions of the rolls are 35 cms length (30 cms working length) and 15 cms diameter. The two-roll mill is run at a temperature of 190°C and a speed of 26/20 rpm (front/rear rolls). Once the film has formed on the front roll and
20 after the standard practice of obtaining a homogenous film, the mill is stopped and a horizontal strip, approximately 4 - 5 cms wide, is cut at the full length of the roll, taken from the roll and stretched immediately to twice its length
25 using a guide of the appropriate length. The strip is held in the stretched position until cool.

The milled-stretched strips from each compound are rated for the amount of undispersed processing aid particles. This is done by comparing gel colonies with
30 reference standards using a rating system of 0 to 100 units, with 0 being equal to a strip having no undispersed particles and 100 being equal to a strip having a large number of undispersed particles. A dispersibility rating of between 40 and 50 units is termed as very good, whereas

anything above 65 - 70 units has an unacceptable number of undispersed gels.

In each series of dispersion evaluations, at least one sample of PVC containing a processing aid without a dispersion aid additive is run as a control.

In order to determine whether a given candidate dispersing aid should be regarded as effective to substantially reduce gel colonies of processing aid in rigid thermoplastic matrix polymer formulations, and is therefore within the scope of this invention, the maximum amount of dispersing aid, (a) to (b) ratio of 98:2, is compared with the control and a reduction of at least 30 units is required. Preferred dispersing aids give a reduction of 45 units and most preferred a reduction of 50 units, ideally 60 units when compared to a processing aid in rigid thermoplastic giving a dispersion rating of 100 (worst).

In the following Examples, the following abbreviations are used:

Methyl methacrylate	= MMA
Butyl methacrylate	= BMA
Styrene	= St
Methyl acrylate	= MA
Ethyl acrylate	= EA
Butyl acrylate	= BA
Butadiene	= Bd

Unless otherwise indicated, all of the candidate dispersing aids in the Examples pass the test set out above.

EXAMPLES 1 TO 8

In these Examples, reported in Table I, various low molecular weight dispersion aids are prepared in emulsion using 4% t-DDM in the above described procedure. The molecular weights of the dispersion aids ranged from 19,000 to 38,000, depending on monomer used. The emulsion is mixed

with an emulsion of a methyl methacrylate-ethylacrylate polymer. The improved processing aids containing the dispersion aids were isolated by spray drying. The dispersion aid is incorporated at a concentration of 3% based on processing aid composition. The ratio of improved processing aid composition (a) and (b) to PVC is 2 to 100. A control is run without the dispersing aid (Example 8).

TABLE I

10	EXAMPLE	DISPERSING AID POLYMER (b) OF	DISPERSION RATING	
			3% based on PVC RATING	% CHANGE
				(relative to control 100 units)
	1	(MMA) (Comparative)	90	10
15	2	(EA)	65	35
	3	(St) (Comparative)	85	15
	4	(BA)	55	45
	5	BA/St : 55/45	40	60
	6	MMA/EA : 90/10 (Comparative)	75	25
20	7	Bd/St : 70/30	60	40
	8	Processing Aid Control (Comparative)	100	-

EXAMPLES 9 TO 13

In these Examples, the relationship between polymer compositions and molecular weight is explored. A series of

low and high molecular weight dispersing aids was prepared by using 4% and 0% t-dodecyl mercaptan chain transfer agent respectively. The molecular weights of the low molecular weight dispersion aids ranged from 19,000 to 38,000, whereas the high molecular weight dispersion aids ranged from 137,000 to 3,890,000. Dispersion data was obtained in the procedure of the previous Examples and are presented in Table II.

Poly-BA and Poly-EA are effective dispersing aids over the range of molecular weights studied. The effectiveness of the dispersing aids seems to diminish very slightly with decreasing molecular weight but not to the point that they are no longer regarded as effective dispersing aids. Similarly, Poly-MMA, Poly-St do not appear to be effective dispersing aids for processing aids at high or low molecular weights.

TABLE II

EFFECT OF DISPERSING AID MOLECULAR WEIGHT
ON THE DISPERSION OF THE PROCESSING AID MODIFIER IN PVC

EXAMPLE	POLYMER COMPOSITION	<u>4% t-DDM</u>		<u>No RSH</u>	
		<u>RATING</u>	<u>3% ADDITIVE ON MODIFIER</u> <u>% CHANGE</u>	<u>RATING</u>	<u>% CHANGE</u>
9	(MMA)				
	(Comparative)	90	10	95	5
10	(EA)	65	35	50	50
11	(St)				
	(Comparative)	85	15	90	10
12	(BA)	55	45	50	50
13	Processing Aid				
	Control				
	(Comparative)	100	-	100	-

EXAMPLES 14 TO 19

The dispersing aids from Examples 2 and 5 were examined for the effect of the additive level on the dispersion of the processing aid in PVC.

5 The sample evaluated in Example 14 showed 44% improvement (compared to Example 15 control) in processing aid dispersion at only 1.5% additive level which is equivalent to only 0.15% on total PVC. In other words it is equivalent to only 0.15% of dispersing aid (a) for every one
10 part of processing aid (b) added to 100 parts of PVC e.g. if 3 phr of (b) is added this will equal 0.45% of dispersing aid (a) for 100 parts of PVC. The results are reported in Table III.

15 The samples evaluated in Examples 17 and 18 show improvement in processing aid dispersion at 2.5% to 5.0% additive level even when mixed with a processing aid (Example 19) which has unacceptably high dispersion rating, being above 100. The dispersing aid is mixed with the processing aid (in emulsion form) in every case in Table III
20 except Example 15 (control) which is a processing aid without any dispersing aid. Example 16 shows the addition of p-MMA as a dispersing aid mixed in emulsion form to a processing aid.

TABLE III

EFFECT OF ADDITIVE LEVEL ON THE
DISPERSION OF THE PROCESSING AID IN PVC

5	EXAMPLE.	DISPERSING AID COMPOSITION	DISPERSION RATING			
			ADD. AT 1.5% ¹		ADD. AT 3% ¹	
			RATING	% CHANGE ²	RATING	% CHANGE ²
	14	Ba/St	50	44.0	40	56
	15	Processing Aid				
		Control				
10		(Comparative	90	-	90	-
	16	MMA (Comparative)	90	-	90	-
			ADD. AT 2.5% ¹		ADD. AT 5% ¹	
			RATING	% CHANGE ²	RATING	% CHANGE ²
	17	(EA)	85	15	55	45
15	18	(Ba/St)	90	10	65	35
	19	Processing Aid				
		Control				
		(Comparative)	>100	-	>100	-

FOOT NOTES

- 20 ¹ the percentages refer to the addition of the dispersing aid to the processing aid,
- ² the percentage change is always compared to the control i.e. Examples 15 or 19.

CLAIMS

1. The use of at least one homopolymer and/or copolymer containing at least 50% by weight of units of one or more of the following monomers: (C₁-C₈)-alkyl acrylates, (C₁-C₉)-cycloalkyl acrylates, ethylene, propylene, vinyl acetate and butadiene as a dispersion aid in a processing aid composition useful in vinyl chloride polymer compositions.
2. A processing aid composition comprising a blend of (a) at least one copolymer of methyl methacrylate and, in an amount of less than 50% by weight of the copolymer, (C₁-C₄)-alkyl acrylate(s) and (b) at least one homopolymer and/or copolymer containing at least 50% by weight of units of one or more of the following monomers: (C₁-C₈)-alkyl acrylates, (C₁-C₉)-cycloalkyl acrylates, ethylene, propylene, vinyl acetate and butadiene; wherein the weight ratio of (a):(b) is 99.5:0.5 to 90:10.
3. Composition as claimed in claim 2 wherein the weight ratio of (a):(b) is 99:1 to 95:5.
4. Composition as claimed in claim 2 or 3 wherein component (b) comprises at least one of the following polymers:
poly(butyl acrylate), poly(ethyl acrylate), polybutadiene, poly(ethyl acrylate/styrene), poly(butyl acrylate/styrene), poly(butyl acrylate/methyl methacrylate), poly(ethyl acrylate/butyl methacrylate), poly(butadiene/styrene), poly(butyl acrylate/styrene/methylmethacrylate) with the proviso that the methacrylate and/or styrene content is less than 50% by weight in each case or poly(vinyl acetate) or poly(ethylene/vinyl acetate).
5. Composition as claimed in any of claims 2 to 4 wherein the molecular weight of component (b) is 15,000 to 100,000 weight average.

6. A composition as claimed in claim 4 or 5 wherein component (b) comprises poly(butyl acrylate), poly(ethyl acrylate) or poly(butyl acrylate/styrene).

7. A composition as claimed in any of claims 2 to 6 wherein component (b) is crosslinked and/or graft linked preferably to a maximum content of 5% by weight crosslinker or graft-linker units.

8. A composition as claimed in any of claims 2 to 7 wherein component (a) comprises poly(ethyl acrylate).

9. A process for preparing a composition as claimed in any of claims 2 to 8 comprising intimately mixing components (a) and (b).

10. A process as claimed in claim 9 wherein component (b) is formed as a latex by emulsion polymerisation in the presence of component (a) as a latex polymer.

11. A process as claimed in claim 9 wherein separate emulsions of components (a) and (b) are prepared and then intimately mixed.

12. A process as claimed in claim 10 or 11 including the further step of coisolating the mixed components, preferably by spray drying or coagulation.

13. A vinyl chloride polymer composition comprising (i) polymer(s) and/or copolymer(s) of vinyl chloride containing at most 50% by weight of other monomer units and (ii) a processing aid composition as claimed in any of claims 2 to 8, the weight ratio of (i) to (ii) preferably being 90:10 to 99.5:0.5.

14. A polymer composition as claimed in claim 13 wherein the processing aid composition has been prepared by a process as claimed in any of claims 9 to 12.

15. A polymer composition as claimed in claim 13 or 14 wherein component (i) comprises poly(vinyl chloride).



European Patent
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EUROPEAN SEARCH REPORT

0132317

Application number

EP 84 30 4347

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl. ³)
X	EP-A-0 037 686 (ROHM & HAAS) * claims * -----	1-6, 8- 9, 11, 13-15	C 08 L 27/06
			TECHNICAL FIELDS SEARCHED (Int. Cl. ³)
			C 08 L
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 15-10-1984	Examiner FOUQUIER J.P.
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	